What is claimed is:

- A method of non-enzymatic ligation of a nucleic acid, comprising contacting a polynucleotide-3' phosphorothicalte with an acceptor polynucleotide under conditions that allow formation of a phosphodiester bond between said polynucleotide-3' phosphorothicalte and said acceptor polynucleotide.
- The method of claim 1, wherein said polynucleotide-3' phosphorothiolate further comprises a 3'
 SNP moiety.
 - 3. The method of claim 1, wherein said polynucleotide-3' phosphorothiolate further comprises a duplex polynucleotide.
- 4. The method of claim 1, wherein said acceptor polynucleotide further comprises a duplex polynucleotide.
 - 5. The method of claim 1, further comprising transducing into a host cell a polynucleotide-3' phosphorothicate having a phosphodiester bond with said acceptor polynucleotide.

6. The method of claim 1, further comprising the step:

contacting a polynucleotide-3'
phosphorothiolate precursor and an activator under
conditions sufficient to react said polynucleotide-3'
phosphorothiolate precursor and said activator to produce
said polynucleotide-3' phosphorothiolate.

- 7. The method of claim 6, wherein said activator 10 is iodonitrobenzene.
- 8. A method of molecular cloning comprising, contacting an insert comprising a polynucleotide-3' phosphorothicate with an acceptor vector under conditions that allow formation of a phosphodiester bond between said insert and said acceptor vector to generate a vector comprising an insert polynucleotide.
 - 9. The method of claim 8, further comprising transforming said vector comprising an insert polynucleotide into a host cell.

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- 10. The method of claim 8, wherein said polynucleotide-3' phosphorothiolate further comprises a 3' SNP moiety.
- 11. The method of claim 8, further comprising the 25 step:

contacting a polynucleotide-3' phosphorothiolate precursor and iodonitrobenzene under conditions sufficient to react said polynucleotide-3'

phosphorothiolate precursor and said iodonitrobenzene to produce said polynucleotide-3' phosphorothiolate.

- 12. A method of molecular cloning comprising,

 5 contacting a vector comprising a polynucleotide-3'
 phosphorothicate with an acceptor polynucleotide, under
 conditions that allow formation of a phosphodiester bond
 between said vector and said acceptor polynucleotide to
 generate a vector comprising said acceptor polynucleotide.
- 13. The method of claim 12, further comprising transforming said vector comprising said acceptor polynucleotide into a host cell.
- 14. The method of claim 12, wherein said
 15 polynucleotide-3' phosphorothiolate further comprises a 3'
 SNP moiety.
 - 15. The method of claim 12, wherein said vector further comprises a 3' phosphorothiolate moiety at one or more terminal ends.

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16. The method of claim 12, further comprising the step:

contacting a polynucleotide-3'
phosphorothiolate precursor and an activator under
conditions sufficient to react said polynucleotide-3'
phosphorothiolate precursor and said activator to produce
said polynucleotide-3' phosphorothiolate.

17. A kit, comprising:

- (a) a polynucleotide-3'
 phosphorothiolate; and
- (b) a buffer in an aqueous solution.

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- 18. The kit of claim 17, wherein said polynucleotide-3' phosphorothiolate further comprises one or more 3' phosphorothiolate moieties.
- 19. The kit of claim 17, wherein said polynucleotide-3' phosphorothiolate further comprises a single stranded polynucleotide.
- 20. The kit of claim 18, wherein said single stranded polynucleotide further comprises anoligonucleotide.
 - 21. The kit of claim 17, wherein said polynucleotide-3' phosphorothiolate further comprises a duplex polynucleotide.
- 20 22. The kit of claim 17, wherein said polynucleotide-3' phosphorothiolate comprises a 3'-SNP moiety.

23. A kit, comprising:

- (a) a polynucleotide-3'
 phosphorothiolate precursor; and
- (b) an activator.

- 24. The kit of claim 23, wherein said polynucleotide-3' phosphorothiolate further comprises one or more 3' phosphorothiolate moieties.
- 5 25. The kit of claim 24, wherein said polynucleotide-3' phosphorothiolate further comprises a single stranded polynucleotide.
- 26. The kit of claim 24, wherein said single stranded polynucleotide further comprises an10 oligonucleotide.
 - 27. The kit of claim 24, wherein said polynucleotide-3' phosphorothiolate further comprises a duplex polynucleotide.
- 15 28. The kit of claim 24, wherein said polynucleotide-3' phosphorothiolate comprises a 3'-SNP moiety.
- 29. A method of ligating a nucleic acid, comprising contacting a polynucleotide-5' phosphorothiolate with a non-sequence specific topoisomerase, or a fragment or modification thereof, and an acceptor polynucleotide under conditions that allow formation of a phosphodiester bond between said polynucleotide-5' phosphorothiolate and said acceptor polynucleotide, with the proviso that said polynucleotide-5' phosphorothiolate does not contain the nucleotide sequence G(C/T)CCTT (SEQ ID NO:5).

- 30. The method of claim 29, wherein said topoisomerase is human topoisomerase I, or a fragment or modification thereof.
- 31. The method of claim 30, wherein said human topoisomerase I is Topo65, or a fragment or modification thereof.
 - 32. The method of claim 29, wherein said polynucleotide-5' phosphorothiolate further comprises a duplex polynucleotide.
- 10 33. The method of claim 29, wherein said acceptor polynucleotide further comprises a vector.
 - 34. The method of claim 29, wherein said polynucleotide-5' phosphorothiolate further comprises a vector.
- polynucleotide-5' phosphorothiolate further comprises a polynucleotide having a 5' phosphorothiolate moiety incorporated within four base pairs from a 3' end of said polynucleotide-5' phosphorothiolate.

36. A kit, comprising:

(a) a polynucleotide-5'
phosphorothiolate, with the proviso that said
polynucleotide-5' phosphorothiolate does not contain a
nucleotide sequence selected from the group of
SEQ ID NO:5, SEQ ID NO:6 or SEQ ID NO:7;

(b) a non-sequence specific topoisomerase, or fragment or modification thereof having topoisomerase activity.

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- 37. The kit of claim 36, wherein said topoisomerase is human topoisomerase I, or a fragment or modification thereof.
- 38. The kit of claim 37, wherein said topoisomerase is Topo65, or a fragment or modification thereof.
 - 39. The kit of claim 36, wherein said polynucleotide-5' phosphorothiolate is a single stranded polynucleotide-5' phosphorothiolate.

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- 40. The kit of claim 37, wherein said single stranded polynucleotide-5' phosphorothiolate further comprises an oligonucleotide.
- 25 41. The kit of claim 36, wherein said polynucleotide-5' phosphorothiolate further comprises a duplex polynucleotide-5' phosphorothiolate.

- 42. The kit of claim 41, wherein said duplex polynucleotide-5' phosphorothiolate further comprises one or more terminal end overhangs.
- 43. The kit of claim 42, wherein said one or more terminal end overhangs further comprise a nucleotide sequence complementary to one or more restriction endonuclease cleavage sites.
 - 44. A composition comprising,
- 10 (a) a polynucleotide-5' phosphorothiolate, with the proviso that said polynucleotide-5' phosphorothiolate does not contain a nucleotide sequence selected from the group of SEQ ID NO:5, SEQ ID NO:6 or SEQ ID NO:7; and
- 15 (b) a non-sequence specific topoisomerase, or fragment or modification thereof having topoisomerase activity.
- 45. The composition of claim 44, wherein said topoisomerase is human topoisomerase I, or a fragment or modification thereof.
 - 46. The composition of claim 45, wherein said topoisomerase is Topo65, or a fragment or modification thereof.
- 25 47. The composition of claim 44, wherein said polynucleotide-5' phosphorothiolate is a single stranded polynucleotide-5' phosphorothiolate.

- 48. The composition of claim 47, wherein said single stranded polynucleotide-5' phosphorothiolate further comprises an oligonucleotide.
- 49. The composition of claim 44, wherein said polynucleotide-5' phosphorothiolate further comprises a duplex polynucleotide-5' phosphorothiolate.
- 50. The composition of claim 49, wherein said duplex polynucleotide-5' phosphorothiolate further comprises one or more terminal end overhangs.
 - 51. The composition of claim 50, wherein said one or more terminal end overhangs further comprise a nucleotide sequence complementary to one or more restriction endonuclease cleavage sites.

52. A compound of the formula:

O = P O H O H P O S R 3

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wherein,

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X is a nucleotide;

y is a positive integer;

R1 is a nucleotide base;

R2 is H or OH; and

R3 is a halo, alkyl, substituted alkyl,

sulfonate moiety, phenyl,

substituted phenyl.

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53. The compound of claim 52, wherein

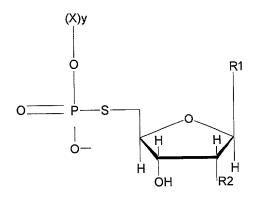
R2 is H.

54. The compound of claim 52, wherein

R3 is nitrophenyl.

55. The compound of claim 52, further comprising a complementary polynucleotide.

56. A compound of the formula:



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wherein,
X is a nucleotide;
y is a positive integer;
R1 is cytosine or guanine; and
R2 is H or OH.